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CHEMICAL ABSTRACTS, vol. 93, no. 22, Decamber 1860, pages 291,292, no. 209118j, Columbus Ohlo (USA);

PHYSICS AND CHEMISTRY OF GLASS, vol. 19, no. 1, February 1978, pages 1-4; A.G. DUNN et al.: "Near infrared optical absorption of iron (ii) in total ecolium borosilicate glasses"

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التعلقم أتمودتا

The present invention relates to a glass suitable for use in the cores of optical fibres, especially for use se core glasses in the production of step-index high numerical sparture (NA) optical fibres, most especially by the double crudible process. The present invention further relates to methods of making such compositions and to optical fibres containing them.

High NA optical fibres are particularly useful for short optical fibre links (e.g. of length less than 1 km) where efficient power leunching is of considerable importance. Among such short links are data links between computers and ancillary equipment and also links in telephone exchanges.

in order to facilitate coupling to other system components such fibres may be required to have large core diameters of for example 175 µm and, moreover, to have an ecceptable level of loss. As the optical fibres are principally intended to be utilised over relatively short links losses of up to 10 dB/m or in some cases more (e.g. up to 15 or 20 dB/km) are acceptable, atthough it is clearly desirable to reduce the loss as much as possible within the constraints.

The numerical aparture of a fibre is theoretically equal to

where \mathbf{n}_1 is the refrective index of the core and \mathbf{n}_2 is the refrective index of the dedding. The amount of light coupled into an optical fibre, other things being equal, is usually proportional to the product of the square of the numerical aperture and the cross-sectional area of the fibre core.

In our European patern application 0018110A we described, inter elia, core glasses suitable for use in optical fibres which had refractive indices in the range from 1,540 to 1,610, were not subject to devitrification or phase separation, and which contained the five components MagO (soda or codium oxide). B_2O_3 (bonic oxide), BaO (baris or barium oxide), SiO_2 (silica or silicon dioxide), and GeO. (germania or germanium dioxida) and only small amounts of other components. Both barium oxide and germanium dioxide are highly refractive components in sodium bereallicate discess, i.e. their presence raises the refractive index of the glass. In order to minimise loss in the eventual fibre. It was necessary to use intermediate reducing conditions in the production of these glasses, these conditions corresponding to a partial pressure of oxygen of about 10⁻⁶ atmospheres. More oxidising etmospheres. More oxidising conditions than this led to increased loss through absorption by transition mazal impurities such as copper while more reducing conditions (e.g. corresponding to a partial oxygen pressure of 10⁻¹³ atmospheres) led to a very high scatter loss men unbown causes.

The $Wa_2O:B_2O_0$ moler ratio in these glasses was high (substantially in excess of 2:1).

The use of these gissess to produce fibres by the double crucible process was also described. In European patent application A—000282, a gisse comprising Na₂O, B₂O₃ BaO (or CaO) and SiO₃, but not ZrO₂, is disclosed. The composition of the glass is, such that it is not subject to devitrification or phase separation during optical than produced on the glass.

of the glass is, such that it is not subject to devitrification or phase separation during optical fibre production. A method of preparing the glass is also disclosed which comprises the steps of preparing a melt (including about 0.1 mole per cent of Al₂O₈ as a redox buffering oxide) and bubbling a mbiture of carbon monoxide and carbon dioxide! through the molten glass.

The use of ZrO₂ (zirconts or zircontum dioxide) as a highly refractive component in vitreous silica for optical fibres is known from our UK parent 1 388 093.

M Yoshiyagawa, Y Kaite, T Ikuma and T Kishimoto, J Non-Crystalline Solide, 40, 489—497 (1990) refers to the use, alone or in combination, of a range of caides in sectium boroellicate places: for optical fibres. These exides are MgO, CaO, SrO, BeO, IZnO, Al₂O₂, La₂O₃, TiO₃, GeO₂ and ZrO₃. No preference among these exides is stated and detailed compositions are not specified.

In Chemical Abstract No. 83:209118] (1990) and Japanese uniscardined patent application 80 60040 there are described sodium borosilicate glasses containing barium code and zirconium dioxide but these have very high barium code contents and very high refractive indexes ranging from 1.69 to 1.65. The same document also discloses glasses containing substantial quantities of further components in combination and not in combination with 8s0 and 2rO₂. Oddas whose uses are suggested and exemplified are U₂O, K₂O, Rb₂O, C₂O, MigO, CeO₇ SrO. Al-O. La-O₂. TiO₂, and GeO₂.

SrO. ZnO. Algo, Legos, Trop, and GeOg. In US patent 4 265 687 (equivalent to French published patent specification 2 481 261) there are described various stable sodium borostillcate glassee containing BaO and ZrOg. These have very high BaO contents, have refrective indexes in excess of 1.81, and contain further components, for example Ligo, K.O., RbgO, Cago, MgO, CaO.

SrO, ZnO, Al₈O₈, L₈O₂, TrO₂ and GeO₂. In French petent application 2 438 018 (equivalent to UK patent application 2 034 300) there are disclosed sodium borosilicate gleases containing BaO and ZrO₂ which have refrective indexes of less than 1.840. There is additionally a disclosure of a glease of refractive index exceeding 1.61 similar to those described in Chemical Abstracts No. 93: 209 118; and already referred to. In the said French patent application, the use is auggested and examplified also of further components in combination with and not in combination with 880 and ZrO₂. Such components are Li₂O, K₃O, Ce₂O, Ti₂O, MgO, CaO, ZnO, PhO, ZnO, Al₂O₃, and TiO₃. The main concern of the French patent application is the production of cladding glasses where water resistance is a prime consideration.

The present invention is based on our surprising discovery that estisfactory glasses

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having refractive indexes in the range 1.540 to Talimis of the town sagamenta gaived furth 018.F glasces of European parent (1018110A) can be produced by the use as components of Na₂O, B_2O_2 , B_2O_3 , B_2O_3 , and E_2O_3 with only minor quantities of other components, the components being used in proportions quite dissimilar from the glasses of the aforcade prior est on BaO- and ZrOz containing sodium borositicate glasses. From the aforesald prior art on 8a0- and ZrO2containing endium korosificate glasses it could not have been predicted that the glasses provided by the present invention would be settlefactory, since it is well known in this field that substantial variations of composition from known satisfactory compositions can affect stability of a glass and the optical tosses in the glass. (In this respect, and generally, it may be noted that glasses which are apparently stable but have ocmpositions close to unstable ones, frequently yield fibres having high loss).

The present invention provides a glass suitable for use in the core of an catical fibre and having a composition such that it is not subject to devitrimention or phase experation, which glass (i) has a refractive index in the range from

1,500 to 1.610 and

(ii) comprises the five components Na₂O, B₂O₂, BaO, SIO, and a highly refractive component other than 8sO and not more than 5 weight per cent of any further components taken together relative to the said five components tetian together, the proportion of 840 being in the range from 2 to 12 mole per cent relative to the said five components taken together and the proportion of SiO₂ being in the range from 40 to 63 mole per cent relative to the said five components taken together,

characterised in that

(i) the said highly refractive component is ZrO₂ in a proportion tying in the range from 1.5 to 18 mole per cent relative to the said five components taken together, the proportion of STO₂ and ZrO₂ taken together being not more than 65 mole per cent relative to the said five components taken together, and

(ii) the Na₂O:B₂O₃ moler ratio is in the range from 1.1:1 to 1.5:1.

An edvantage of these glasses over the aforesald glasses of European patent application 001B110A is that they do not display scatter loss when highly reduced. This parmits the bast possible suppression of absorption loss due to transition metal lons such as copper by a convenient process.

A further advantage is that the amount of ZrO_n required to achieve a given refractive index is leas then the amount of GsO_D so that the purity specification for the ZrO, need not be so stringent for a given meetinum prescribed lavel of interfering impurides. There are additionally overall cost advantages to the use of ZrO₂ of the requisite purity in the requisite quantities. Cost advantages

B are especially important for large diameter cores because of the amount of material involved.

The glaces according to the present invention are generally more viscous when motion than the aforesaid glosses of European patent application 0018110A, which offers interesting new possibilities. In particular, it offers a way of evoiding or reducing the problem that can be encountered with these prior art glasses, namely the excessively rapid pulling of core glass compared with the clading glass in the cauble crucible process

On the various compositional features of the glass according to the present invention further observations may be made as follows, without explicit comparison with the prior art.

The limit of 5 weight per cent on further components reflects the fact that (apart from agents) possibly recor buffering components can usually be avoided. This is good because, firstly, the use of numerous components esta to notizently expensive question of the purity specification with each one and, secondly it simplifies production of the glass. Apart from redox buffering agents, a minor compenent which was presently think may prove attractive is

Al₂O₂, perhaps at a lavel of 1 to 3 weight per cent.
The lower limit on the proportion of ZrO₂ ensures that at least a part of the refrective index increase is due to zirconia. With BaO alone, it is difficult to schieve refractive indices much in excess of 1.55 without the glass becoming unstable. However, even with the use of ZrO₂ a molar proportion in excess of 15 per cent tends to lead to instability (albeit for a much higher refractive index).

The range for moler ratio Na₂O:B₂O₂ Is especially suitable in achieving a good balance between glass stability (favoured by Na_sO) and fibre-forming performance in the double crucible process (favoured by B.O.).

Preferably, the proportion of ZrOs in the present invention is in the range from 3 to 10 mole per cent. Preferably, the proportion of BaO is in the range from 4 to 10 mole per cant.
The moles ratio Ne₃O:B₂O₃ is preferably in the

range 1.10 to 1.50, especially from 1.10 to 1.40.

The present invention affords glasses with especially good viscosity properties for the range of refractive Index 1.550 to 1.590.

Preferably the propurtion of Na₂O lies in the range from 15 to 25 mole per cent and that of $B_{\rm g}O_{\rm d}$ lles in the range from 10 to 20 mole per cent, both proportions being relative to the five components (Na₂O, B₃O₂, 8aO, SiO₂, and ZrO₂) taken together. Preferably, the proportion of SiOs and ZrOs taken together is at least 50 mole per cent relative to the said five components taken together, and more preferably at lesses 65 mole per cent.

It will be exprecised that the glass according to the present invention can be formed in any appropriate manner, and that formation as a mait which can be subsequently converted to a solid glass for storage is especially convenient.

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The present invention further provides a method of preparing the glass provided by the invention which comprises the steps of

(a) preparing a melt including from 0.01 to 1 weight per cent of a redox buffering egent or of one or more redox buffering agants taken together relative to the eald five components taken together, and

(b) passing carbon monoxide through the mait. We find amenic trioxide, As₂O₂, to be a convenient redox buffering agent, but other oxides, for example antimony oxide (Sb₂O₂), can also be used. Preferably, the amount employed lies in the range 0.05 to 0.2 weight per cent. The function of these agents is described in general terms in European petern application 0018110A and in UK patent 1 507 711 referred to therein.

In general, melt composition and final glass composition are practically the same, and any deviations can be allowed for after simple trial and error. We have observed minor deviations due to reduction of As₂O₃ to volatile As during step (b).

Carbon monoxide is conveniently used in the form of a mixture with carbon dioxide, said mixture however preferably not containing less than 10 per cent by volume of carbon monoxide.

Depending on the transition metal contaminents in the malt, it may be desirable to pass oxygen through the malt (either pure or in admixture with other gases) prior to step (b).

The present invention also provides an optical fibre comprising a core comprising the glass provided by the present invention and a cladding comprising a sodium borosilicate glass of lower refractive Index.

The sodium borosilicate glass used for the cladding may include components other than Na₂O, B₂O₂, and SiO₂. Preferably, only small quantities of such further components are used. e.g. Al₂O₂ et a level of 1—2 mole per cent relative to Na₂O, B₂O₃ SiO₂, and Al₂O₃ taken together may afford advantages of durability and water resistance. Preferably, the glass is in a highly reduced state.

Preferred fibres in accordance with the present invention are made by the double crucible process. Convenient large core high NA fibres in accordance with the present invention have a core diameter in the range from 90 to 220 µm. The outer diameter of the dedding is normally at least 25 pm greater than the care diameter and is conveniently in the range from 115 µ to 270 µm.

The present invention will now be further illustrated by means of Examples 1 to 12 and Comparative Examples C1 to C5 (not in accordance with the invention).

Each of these Examples and Comparative Examples illustrates a core glass material. (or en attempted core glass material). The compositions and observed properties are given in Tables 1 and 2 respectively.

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TABLE 1 Giasa compositions

Example _	Mole percentages							
No.	Na₌O	B ₂ O ₃	BaQ	ZrO ₂	SiO ₂			
1	20	14.6	4	2	59.5			
2	20	14.5	4	3.5	58			
8	20	14.5	. 4	5	56.5			
4	20	14.5	4	10	51.6			
5	20	14.5	4	12.5	49 .			
6	20	14.5	4	15	46.5			
C1 .	20	14.5	4	16	45.5			
CZ	20	14.5	4	17.5	44			
C3	20	16	4	17.5	42.5			
7	20	18	4	8	50			
8	20	19	4	15	43			
9	19	16	8	. 3.	95			
10	19	15	.9	5.18	52.84			
11	19	15	. 8	. 7	51			
12	19	15	8	io	48			
	Na _z O	B ₂ O ₉	BaO	GeO ₂	SiOz			
C4	20	8.3	. 11.7	12	48			
C5	20	8.3	11.7	25	35			

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TABLE 2 Glass properties

Example No.	ā cm_s b ļu	ain 10 ⁻⁷ °C ⁻¹	n _D	Tg In ⁴C	SP in °C	Stability
1	2.73	81.9	1.5438	670 .	635	Y68
2	2.76	81	1.5509	574	640	Yes
2	2.79	74.8	1,5577	587	860	Yes
4	2.89	76.8	1,5815	592	670	Yes
5	2.94	75.9	1.5915	592	883	Yes
8	2.99	72.5	1.6028	615	672	Yes
C1	_	_	_	_	_	No
C2		_	_	_		No
C3	_	<u></u>	_	_	_	No
7	2.88	88.6	1.5750	676	845	Yes
8	2.97	82.9	1,6017	5 98	668	Yes
9	2.90	81.1	1.5815	578	620	Yeş
10	2.94	79.7	1.5714	575	645	Yes
11	-2.58	82.05	1.5794	595 ⁻	648	Yes -
12	8,05	78.1	1.5936	593	643	Yes
C4	2.11	-	1.6688	523	585	Yes
C15	9.34		1,5916	505	546	Yes

p=density, q=linear coefficient of expansion, n_p=refractive index (NaD line), Tg=glass temperature, SP=diatometric softening point.

The preparative procedure for Examples 1 to 12 and Comparative Examples C1 to C3 was as follows: The starting materials were high-purity Ns₂CO₅, Ns₂B₄O₇, BsCO₉, ZrO₂, SiO₂, and As₂O₃ powders. The powders were mixed together in the proportions appropriate to the desired glass composition, the Ac₂O₃ being used at a level of 0.1 weight per cent relative to the other components taken together. The mixture was then introduced gradually into a silica crucible at 1100°C where it melted. The melt was left to react for a further hour or so end the temperature was relead to 1160 to 1200°C and pure dry carbon dioxide was passed over the top of the melt at 2 litre/min, this flow being maintained to the end of the preparation. Gassa were bubbled through the melt in the following order: pure dry carbon dipoide for 2 hours; pure dry oxygen for } hour; and a mixtue of 18 volume per cent carbon monoxide and 82 volume per cent carbon dioxide for 13 hours. In each case the flow rate was 500 ml/min. Bubbling was then discontinued (the

flow of carbon dicadds over the melt being maintained) and the melt was heated to 1250°C for 18 hours to remove bubbles. The melt was then cooled to 850-900°C, and the glass was removed as rods and stored.

The preparative procedure for Comparative Examples C4 and C5 was in accordance with the invention of European patent epplication 0018110A end in particular the glass was in an intermediate redox state (corresponding to a pertial oxygen pressure of about 10⁻⁸ atmospheres) schleved with the use of pure dry carbon dioxide instead of the carbon monoxide/carbon dicadde mbdure (which latter corresponded to a partial oxygen pressure of about 10-13 atmoepherez).

The three series of Examples 1 to 8, 7 to 8, and 9 to 12 respectively show how, if the molar proportions of Na₂O, B₂O₂, and BaO are fixed, the refractive index varies as one changes the relative amounts of ZrO2 and StO2 constituting the balance of the composition.

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The glass of Example 12 was found to corrode the silica crucible in which it was being meted, but this problem is one which may be overcome by the use of an alternative material for the

Comparative Examples C1 to C3 Illustrate the instability (due to crystallization) that erises for high ZrO₂ concentrations. Comparative Examples C4 and C5 illustrate the lower glass and softening temperatures (and hence viscosities) of the prior art glasses in accordance with European petent application 0018110A.

From glasses in accordance with the present invention, fibres were prepared having core diameters of 175 µm and other disdding diameters of 200 µm. The cladding glassess were prepared with the use of As₂O₃ as a redox buffering agent and with the use of a carbon monoxida/carbon dioxide mixture to achieve a highly reduced state.

An optical fibre was prepared from the glass of Example 2 by the double crudble process. The example 2 by the double crucion process. The drawing (i.e. nowle) temperature was approximately 850°C. The dedding glass had the composition 20 mole % Ne₃O, 17.5 mole % B_2O_3 , 81 mole % SIO₂, and 1.5 mole % Al_2O_3 and had a refractive locate n_0 of 1.5185. The numerical apparture of the fibre was 0.32 and the fibre had a content of 7.3 content of 5.3. loss of 7.3 (13/km at 850 nm.

An optical fibre was prepared also from the glass of Example 7. The drawing temperature was In this case about 900°C because of the higher viscosity of the glass. The cladding glass had the composition 20 mole % Na₂O, 17.5 mole % B₂O₃, 58 mole % SiO₃, 2 mole % NgO, and 1.5 mole % Al₂O₃ and a refractive index of 1.5183. The numerical aparture was 0.41, and the faire had a loss at 850 nm of 10.9 dB/km.

Chims

1. A glass suitable for use in the core of an optical fibre and having a composition such that it is not subject to devitrification or phase separation, which glass

(I) has a refractive index in the range from 1.540

to 1,610 and

(ii) comprises the five components Na₂O, B_2O_2 BaO, SiO₂, and a highly refractive component other than BaO and not more than 6 weight par cant of any further components taken together relative to the said five components taken relative to the said five components taken together, the proportion of BaO being in the range from 2 to 12 male per cent relative to the said five components taken together and the proportion of SiO_2 being in the range from 40 to 63 male per cent relative to the said five components taken

characterised in that (I) the sold highly refractive component is ZrO2 in a proportion lying in the range from 1.5 to 15 male per cant relative to the said five components teiten together, the proportion of SiO_a and ZrO₂ taken together being not more than 65 mote per

10 cent relative to the said five components taken . together, and |

(ii) the Na₂O:B₂O₃ moder ratio is in the range from 1.1:1 to 1.5:1

2. A glass according to claim 1 wherein the proportion of ZrO2 is in the range from 3 to 10 mole per cent

3. A glass according to claim 1 or claim 2, wherein the proportion of BsO is in the range

from 4 to 10 mole per cent.

4. A gless according to any preceding claim, wherein the molar ratio Na₂O:S₈O₃ is in the range from 1.10 to 1.40.

5. A glass according to any preceding claim, wherein the proportion of Na₂O is in the range from 15 to 25 mole per cent reletive to the said

five components taken together.

6. A glass according to any preceding claim, wherein the proportion of B_2O_3 is in the range from 10 to 20 mole per cent relative to the said five components taken together.

7. A glace according to any preceding claim, wherein the proportion of SiO₂ and ZrO₂ taken together is in the range from 50 to 63 mole par cent relative to the said five components taken together.

8. A glass eccording to claim 7, wherein the proportion of SiO₂ and ZrO₂ taken together is in the range from 65 to 65 mole per cent relative to the said five components taken together.

9. A glass according to any praceding claim which has a refrective index in the range from 1,650 to 1,580.

10. A glass according to any precading claim which is in a highly reduced state.

11. A method of preparing a glass according to any preceding claim which comprises the steps of (a) proparing a melt including from 0.01 to 1 weight per cent of a redox buffering agent or of one or more redox buffering agants taken together relative to the said five components tation together, and

(b) passing carbon monoxide through the melt. 12. A method according to claim 11, wherein the redox buffering agent is A9203-

13. A method according to claim 11 or claim 12, wherein the melt of (a) includes from 0.05 to 0.2 weight per cent of a radox buffering agent.

14. A method according to any of clasms 11 to 13, wherein step (b) is performed with a mixture of carbon monoxide and carbon dloxide containing at less: 10 mole per cent of the former.

16. A method according to any of claims 11 to 14, wherein copygen is passed through the malt between step (a) and step (b). 16. An optical fibre which comprises a core

comprising a glass according to any of claims 1 to 10 or prepared according to any of claims 11 to 15

and a sodium borosilicate cladding glass. 17. An optical fibre according to claim 16, wherein the core dismeter is in the range from 80

to 220 µm.i 18. An optical fibre according to claim 16 or cialm 17, which has a loss at 850 nm of not more

than 20 dB/km.

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19. An optical fibre according to claim 18, which has a loss at 850 nm of not more than 15 dB/km.

Petentansprüche

1. Zur Verwendung im Kern einer optischen Faser geeignetes Glas mit einer solchen Zuaammensetzung, daß es keine Emglasung oder Phasentrannung erleidet, welches Glas

(i) einen Brechungsindex im Bereich von 1,540

bis 1,610 hat und

(II) die fünf Bestandteile Ne₂O, B₂O₂, BeO, SiO₂ and einen hochgredig brechenden Bestandteil außer BaO und nicht mehr als (nsgesamt 5 Gew.% irgendweicher weiterer Bestandteile, bezogen auf die Gesamtmenge der fünf Bestandteile, aufweist, wobei der BaO-Anteil im Bereich von 2 bis 12 Mol-%, bezogen auf die Gesamtheit der fünf Bestandteile, liegt und der SiO_x-Anteil im Bereich von 40 bis 63 Mol-%, bezogen auf die Gesamtheit der fünf Bestandteile liegt.

dadurch gekennzeichnet, daß

- (I) der hochgradig brechende Bestandteil ZrO₂ în einem Anteil ist, der Im Bereich von 1,5 bis 15 Mol-%, bezogen auf die Gesamtheit der fünf Bestandseile, liegt, wobsi der Anteil von SiO₂ und ZrO₂ zusammen nicht mehr als 65 Mol-%, bezogen auf die Gesamthelt der fünf Bestandtelle,
- (iii) das Na₂O:8₂O₃-Molverhältnis im Bereich von 1,1:1 bis 1,5:1 lst.

2, Gles nach Anspruch 1, worin der ZrO₂ Antell im Bereich von 3 bis 10 Mol-% ist.

- 3. Glas nach Anspruch 1 oder Anspruch 2, worin de BaO-Antail im Bereich von 4 bis 10 Moi-% Ist.
- 4. Glas nach irgendeinem vorstehenden Anspruch, worm das Moi-verhähnts Na₂O:B₂O₃ im Bersich von 1,70 bis 1,40 ist.
- 5. Glas nach irgandeinem vorstehenden Anspruch, worm der Na₂O-Antell im Bereich von 15 bla 25 Mol-%, bezogen auf die Gasamtheit der
- fünf Bestentbelle, ist. 6. Glas nach Irgendelnem vorstehendes Anspruch, worin der B₂O₂-Anteil im Bereich von 10 bis 20 Mol-%, bezogen auf die Gesamtheit der funf Bestandteile, Ist.
- 7. Glas nach ingandeinem vorstehenden Anspruch, worth der Antell von SIO₂ und ZrO₂ zusämmen im Bereich von 60 bis 65 Mol-%, bezogen auf die Gesamtheft der fünf Bestandteile,
- 8. Gles nach Anspruch 7, worin der Anteil von SiO₂ and ZrO₃ zusammen im Bereich von 55 bis 85 Mol-%, bezogen auf die Gesamtheit der fünf Bestandteile, ist.
- 9. Glas nach irgendeinem vorstehenden Anspruch, das einen Brechungsindex im Bereich von 1,560 bis 1,590 hat.
- 10. Glas nach irgendeinem vorstahenden Anspruch, das in einem hochgradig reduzierten Zustand ist.
- 11. Verfahren zum Herstellen eines Glases nach Irgendelnem vorstehenden Anspruch, das die Schritte des

a) Horstellens einer Schmalze, die 0,01 bis 1 Gew.% eines Radcopuffermittels oder eines oder mehrere Redexpuffermittal insgesemt, bezogen auf die Gesammenge der fünf Bastandhalle, enthält, und

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b) Durchleitens von Kohlenmancodd durch die

Schmelze umfaßt.

12. Verfehren nach Anapruch 11, wobei das

Redoxpuffermittel As₂O₃ lst. 13. Verfehren nach Anspruch 11 oder Anspruch 12, wobei die Schmelze van (e) 0,05 bis 0,2 Gew.% eines Redoxpuffermittels enthält.

14. Verfahren nach ingendalnem der Anaprüche 11 ble 13, wobei der Schritt (b) mit einer Mischung von Kohlenmonoxid und Kohlendioxid durchgeführt wird, die wenigstens 10 Mol-% des ersteren enthält.

15. Verfahren nach irgandeinem der Ansprüche 11 bis 14, wobel zwischen dem Schritt (a) und dem Schritt (b) Sauerstoff durch die Schmelze

geleitet wird.

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16. Optische Faser, die einen Kern mit einem Glas nach ingendeinem der Ansprüche 1 bis 10 oder das nach irgendelnem der Ansprüche 11 bis 15 hergestellt ist, und ein Natriumboreilikatüberzugagias aufweist

17. Optische Hasier nach Anspruch 16, worin der Kerndurchmasser Im Bereich von 90 bis 220 µm

18. Optische Faser nach Anspruch 16 oder Anspruch 17, das einen Verlust bei 860 mm von nicht mehr eis 20 dB/hat.

19. Optische Faser nach Anspruch 16, das einen Verlust bei 850 nm von nicht mehr els 15 dB/km

Revendisations

1. Un verre adapté à la constitution du cosur d'une fibre aptique et sysnt une composition telle qu'il ne soit pas sujet è dévitrification ou séparation de phase, verre qui

(I) a un indice de réfraction comprie entre 1,540

et 1,810 et

(II) comprend les cinq composants Na₂O, B₂O₃, BaO, SIO₂, et un composant à haut pouvoir refringent autre que BaO et pas plus de 5% an poids de tous autres composants pris dans laur ensemble relativement aux cinq dite composante pris dans leuri ensemble, la proportion de BeO étant comprise entre 2 et 12 moies % relativement aux cinq dits composants pris dans leur ensemble et la proportion de SiO2 étant compris entre 40 et 83 molas % rélativement aux conq d'its composants pris dans leur ensemble,

caractérias en ce que

(i) ledit composent hautement refringent est ZrO₂ dans une proportion comprise entre 1,5 et 15 moles pour cent relativement aux cing dits composants pris dans leur ensemble, la proportion de SiO2 et ZrO2 pris dans leur ensemble n'étant pas plus grande que 65 moise pour cent relativement sux cinq dits composants pris dans laur ensemble et

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(II) le rapport molaire Na₂Q:8₂Q₃ est compris entre 1,1:1 et 1,5:1.

2. Un verre selon la revendication 1, caractérisé en ce que la proportion de ZrO₂ est comprise entre 3 à 10 moles pour cent.

3. Un verre selon la ravendication 1 ou 2, caractérisé en ce que le BaO est compris entre 4 et 10 moles pour cent.

 Un verre salon l'une quelconque des revendications précédentes, caractérisé en ce que le repport moleire Ne₂O:B₂O₃ est compris entre 1,10 et 1,40.

5. Un verre selon l'une quelconque des revendications précédentes, caractérisé en ce que la proportion de Na₂O est comprise entre 15 et 25 moles pour cent relativement aux cinq dits composents pris dans leur ensemble.

9. Un verre colon l'una qualconque des revendications précédantes, caractérisé en ce que la proportion de 8₂O₃ est comprise entre 10 et 20 moles pour cent relativement aux cinq dits composants pris dans leur ensemble.

7. Un verre selon l'une quelconque des revendications précédentes, caractérisé en ce que la proportion de SiO₂ et ZrO₂ pris dans leur ensemble est comprise entre 50 et 65 moles pour cent relativement aux cinq dits composants pris dans leur ensemble.

8. Un varre selon la revendication 7, caractérisé en ce que la proportion de SIO₂ et ZrO₂ pris ensemble est comprise entre 55 et 65 moles pour cent relativement aux sing dite composants pris dans leur ensemble.

9. Un verra celon l'una quelconque des revendications précédentes, ayant un indice de réfraction compris entre 1,550 et 1,590.

 Un verre salon l'une quelconque des revendications précédentes, qui est dans un état hautement réduit.

11. Une méthode de préparation d'une varre selon l'une quelconque des revendications précédentes qui comprend les étapes sulvantes

(a) préparation d'un bain de fusion comprenant antre 0,01 à 1% en polds d'un agent jouant le rôle de tampon dans la réaction d'oxydo-réduction ou da l'un ou plusieurs agents jouant le rôte de tampon dans la réaction d'oxydo-réduction pris dans leur ensemble relativement aux cinq dits composants pris dans leur ensemble, et

(b) circulation de monoxyde de carbone au travers dudit bein de fusion.

12. Une méthode salon la revendication 11, caractérisé en ce que l'agent jouant le rôle de tempon dans la réaction d'oxydo-réduction est AS₂O₃.

AS₂O₃.

13. Una méthode selon la revendication 11 ou 12, caractérisé en ce que la bain de fusion de (a) comprend de 0.05 à 0.2 % en poids le l'agent jouant le rôle de tempon dans la réaction d'oxydo-réduction.

14. Une méthode selon l'une des revendications 11 à 13, ceractéries en ce que l'étape (b) est réalisée en utilisant un métange de monocyde de carbone et de dioxyde de carbone contenant au moins 10 moies % de monoxyde de carbone.

16. Una méthode selon l'une qualconque des revendications 11 à 14, caractérisé en ca que de l'oxygène est envoyé à travers le bain en fusion entre l'étape (a) et l'étape (b).

16. Une fibre optique qui comprend un coeur contenant un verre, caractérisé par l'une quel-conque des revendications 1 à 10 ou préparé suivant l'une quelconque des revendications 11 à 16 et une anveloppe de verre an borosilicate de sodium.

17. Une fibre optique selon la revendication 16, caractéries en ce que le diamètre de coeur est compris entre 50 et 220 µm.

18. Una fibre optique salon la revendication 16 ou 17, caractérisé en ca qu'elle a des pertes à 850 nm inférieures à 20 dB/km.

19. Une fibre optique selon la revendication 18, caractérisé en ce qu'elle a des pertes à 850 nm inférieures à 15 dB/km.

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